

Myriad Genetics Corporate Presentation

01/08/2018

Forward Looking Statements

Some of the information presented here today may contain projections or other forward-looking statements regarding future events or the future financial performance of the Company. These statements are based on management's current expectations and the actual events or results may differ materially and adversely from these expectations. We refer you to the documents the Company files from time to time with the Securities and Exchange Commission, specifically, the Company's annual reports on Form 10-K, its quarterly reports on Form 10-Q, and its current reports on Form 8-K. These documents identify important risk factors that could cause the actual results to differ materially from those contained in the Company's projections or forward-looking statements



Our vision...

A trusted advisor transforming patients' lives worldwide with pioneering molecular diagnostics



Myriad Leads Personalized Medicine Industry

Unique Scale and Expertise Creates Sustainable Competitive Advantage

- Over **27** years since company was founded in 1991
- 1 only profitable R&D driven personalized medicine company
- >3.0 million tests performed; unmatched reputation for quality
- **2,500** employees dedicated to the Myriad mission
- >1,000 scientific publications to date; extensive research capabilities
- 3 expertise with all three types of biomarkers (DNA, RNA, and proteins)
- 1 only personalized medicine company with broad regulatory experience
- >90,000 ordering physicians since inception; deep relationships
 - 95% of U.S. payers are in-network



Answering patients' four most pressing questions

Will I get a disease?

Do I have a disease?

Should I treat this disease?

How should I treat this disease?





In **six** medical specialties

Oncology Dermatology

Preventive Care Autoimmune

Urology Neuroscience



Critical Success Factors to Achieving Strategic Goals

STRATEGIC GOALS

CRITICAL SUCCESS FACTORS

>10%

Revenue Growth

>30%
Operating Margin

7 Products

>\$50M

>10%
International
Revenue

Stabilize hereditary cancer revenue

Grow new product volume

Expand reimbursement coverage for new products

Increase RNA kit revenue internationally

Improve profitability with Elevate 2020



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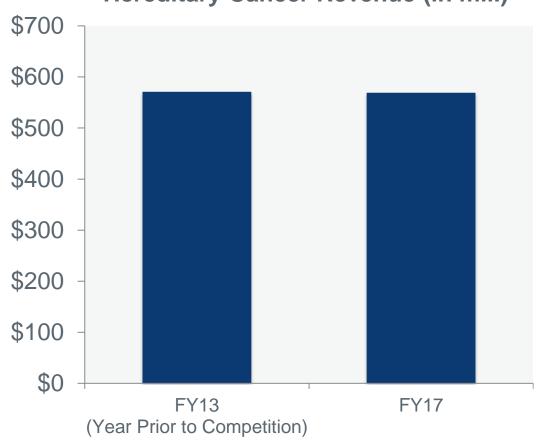
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Hereditary Cancer Business Has Been Durable Significant Competitive Advantages Results in Continued Leadership

Hereditary Cancer Revenue (in mil.)



VOLUME



- Up 15% with flat revenue after 4 years of competition
- > 3% volume growth target in 1Q18
- Third quarter in a row with YoY volume growth

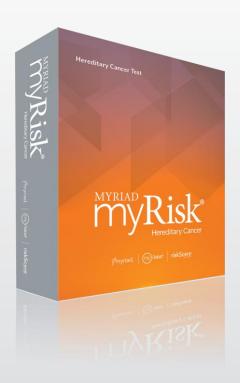
PRICING \$



- 86% under long-term, fixedprice contracts
- Predictable pricing

A Market that Demands Perfection

Three Competitive Advantages Provide Important Differentiation





Lab accuracy:

- 85,000 base pairs with 100% accuracy
- 856 steps using 23 major technology platforms
- 100 proprietary software applications

Variant classification:



- 20 years of research; > \$100M investment
- >2.5M patients tested; 50k variants identified
- 5 proprietary methods with 99.5% validity

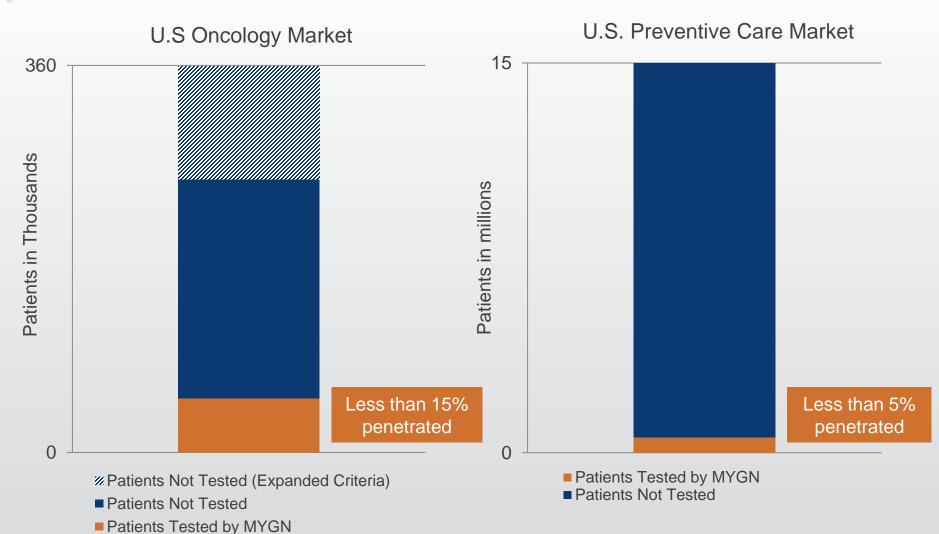
Customer service & support:



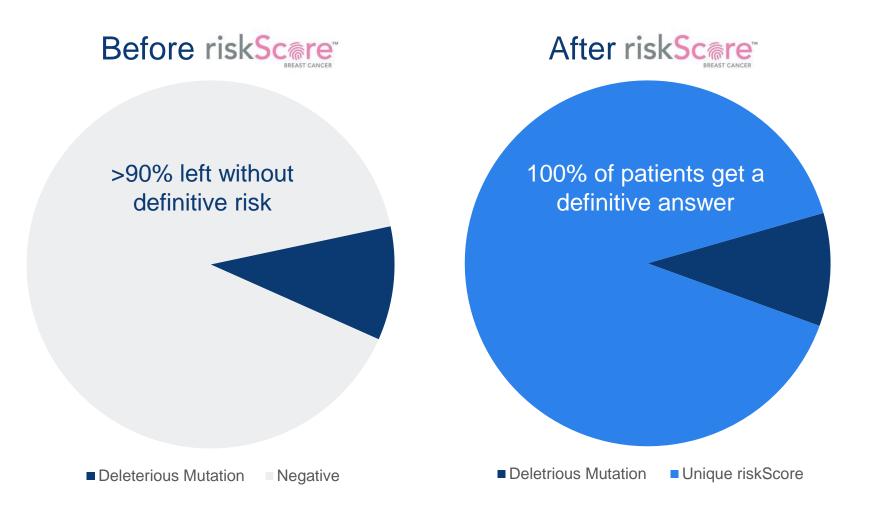
- Over 40k ordering physicians annually
- 450 field educators
- Extensive reimbursement support
- Lifetime commitment to patients

Substantial Opportunity for Volume Growth

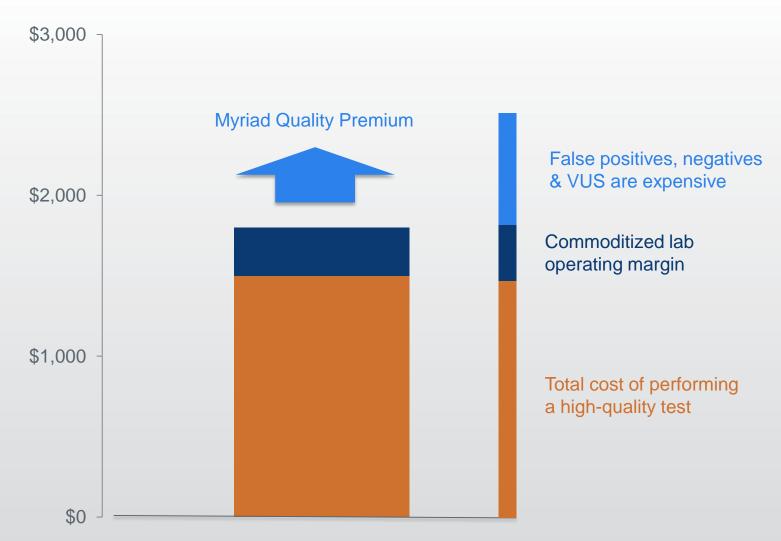
Highly Underpenetrated in Both Oncology and Preventive Care



riskScore Represents New Epoch in Hereditary Cancer Broadens Already Significant Competitive Moat



Cost of Performing High-Quality Test Establishes Pricing Floor Myriad Quality Justifies Significant Price Premium





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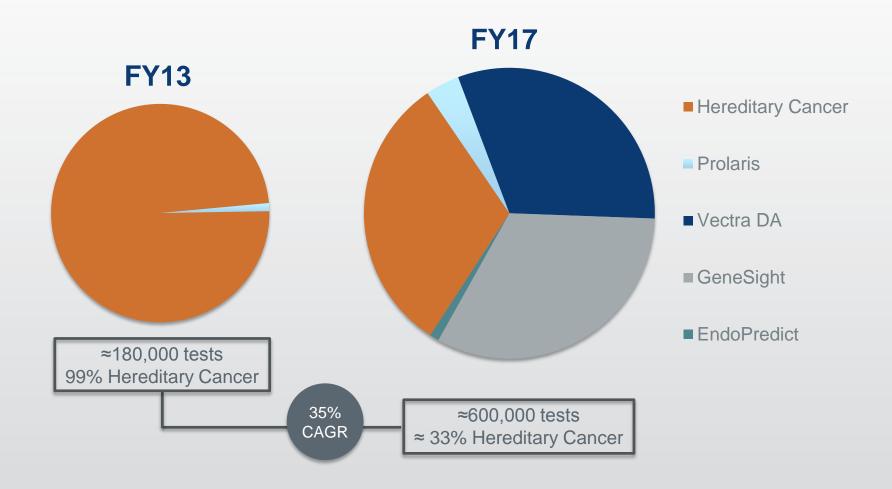
Industry Leading Pipeline

Nine Proprietary Products Sold Through Multiple Channels

Product	Clinical Question	Global Market Size	Technology	Distribution Channel
MYRIAD Risk® Hereditary Cancer riskSc@re™ BREAST CANCER	Will I get cancer?	\$4B 1.5 million patients	28 gene DNA sequencing >80 SNPs + clinical data	Oncology Preventive Care Urology
genesight [*]	What antidepressant should I take?	\$10B 5 million patients	12 genes multiplex PCR proprietary algorithm	Neuroscience Preventive Care
Vectra DA	Should I change my RA treatment?	\$3B 3 million patients	12 protein markers proprietary algorithm	Autoimmune
Prolaris	How aggressively should I treat my prostate cancer?	\$1.2B 400,000 patients	46 RNA markers proprietary algorithm	Urology
EndoPredict®	How aggressively should I treat my breast cancer?	\$0.9B 350,000 patients	12 RNA markers proprietary algorithm	Oncology
MYRIAD Path	Is this skin lesion melanoma?	\$0.8B 600,000 patients	23 RNA markers proprietary algorithm	Dermatology
BRACAnalysis CDx°	Should I use a PARP inhibitor to treat my cancer?	\$6.0B 1.5 million patients	2 gene DNA sequencing tumor DNA sequencing proprietary algorithm	Oncology

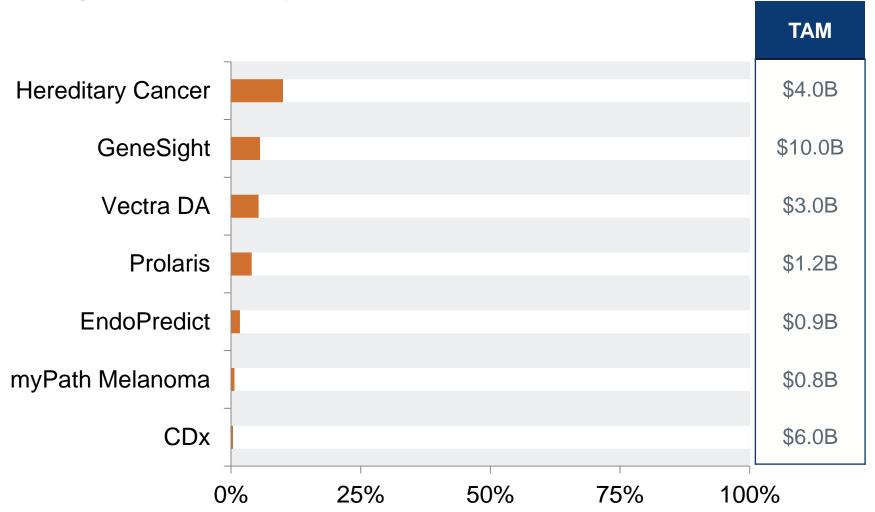
Substantial Diversification in Testing Volumes

≈ Two-Thirds of Volume Generated by New Products



Early Stages of Market Adoption

Significant Opportunity For Continued Growth





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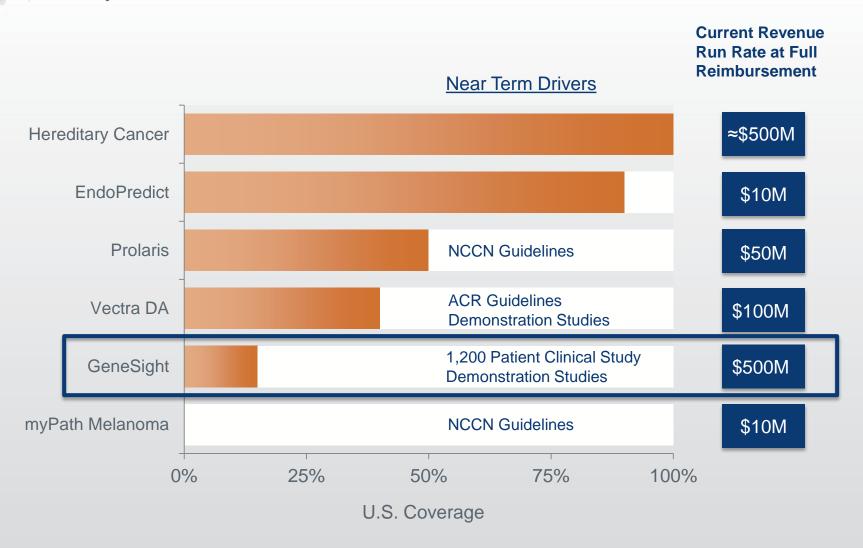
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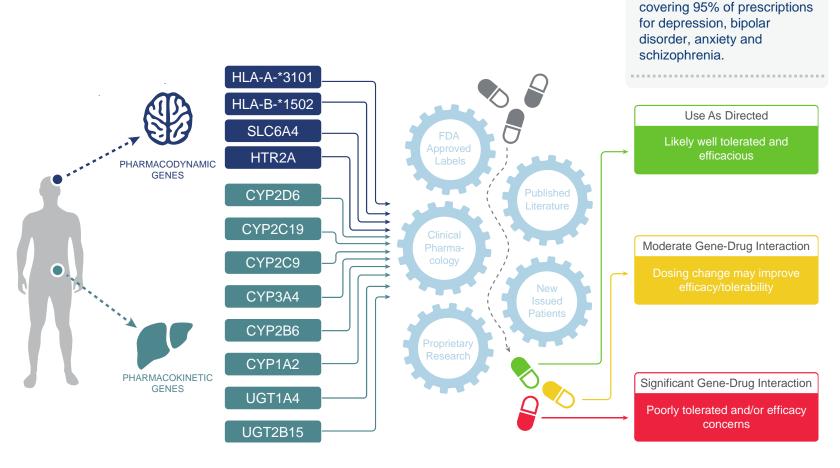
Improve profitability with Elevate 2020

Additional Reimbursement is a Growth Multiplier

Fully Reimbursed New Product Revenue is \$670 Million



GeneSight® Psychotropic Proprietary Combinatorial Pharmacogenomic Test



Data on 55 antidepressants

331,776 unique GeneSight report genetic combinations and medication recommendations

Phase 3 GeneSight RCT Study – Class A Evidence

Overview

- Randomized, 8 week double-blind, controlled evaluation followed by an openlabel follow-up period of an additional 16 weeks
- Compare the patient response to psychotropic treatments in a GeneSight-guided arm (GS) vs. a Treatment as Usual (TAU) arm

Population

- ~1,200 patients,18 years or older
- Diagnosed with moderate to severe depression
- Inadequate response to at least 1 psychotropic treatment

Arms

- GeneSight GS
- Treatment As Usual TAU (any medication, multiple medication, any doses)

Endpoint

- HAMD-17 Scores compared at baseline to 8 week time point
- 3 calculations: Remission, Response, Symptom Improvement

Investigators











GeneSight Study Significant for Most Important Endpoints Beginning Discussions With Commercial Payers

Study endpoint	What it Means	Study Result	Importance to Clinicians and Payers
Remission hardest to achieve	Patient no longer depressed	Highly statistically significant (p<0.01)	Very important
Response difficult to achieve	Patient feels a lot better	Highly statistically significant (p=0.01)	Very important
Symptom Improvement most likely to achieve	Patient feels somewhat better	Approaching statistical significance (p=0.1)	Meaningful

- Remission, Response, and Symptom Improvement were durable and continued to improve over the 24 week study period
- 40 antidepressant FDA registration studies in the last 20 years:
 - All were compared to placebo, not active drug like GeneSight
 - Only 13% showed statistical significance for Remission
 - Only 33% showed statistical significance for Response



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International Growth Focused in Major Geographies Kit Products Appeal to Existing Business Models

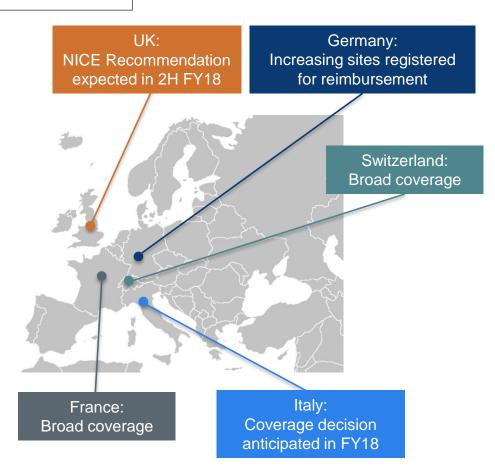
COUNTRIES		KITS		REFERENCE TESTS	
Direct Sales:	EU5 + Canada	RNA (platform partner):	EndoPredictProlarismyPath Melanoma	DNA	myriad Risk ™
Distributor	China, Japan, Brazil, and others	Protein (platform partner):	Vectra DA myPath Bipolar	(multiple platforms):	Companion Diagnostics

EndoPredict Well Positioned For Continued Growth

Major Reimbursement Catalysts Should Drive Market Traction

Focus - EU5 & Canada







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Launching Elevate 2020

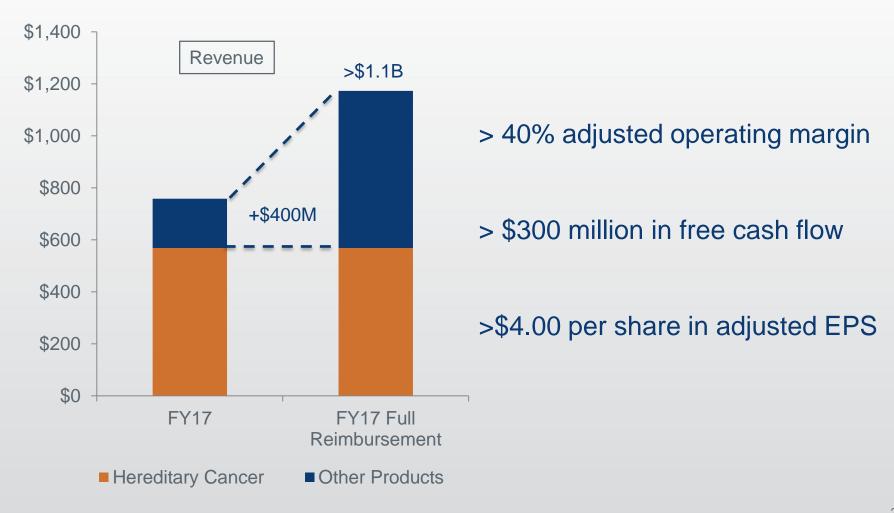
Goal of Achieving \$50M in Incremental Operating Profit by FY20



Financial Outlook

Additional Reimbursement Transformative

With Full Reimbursement FY17 Revenue >\$1.1B, Operating Margins >40%



Potential Catalysts in Fiscal Year 2018 Multiple Possibilities for Material Upsides

Product	Potential Catalyst	Potential Timing	Progress
MYRIAD Risk® Hereditary Cancer	Better Volume Growth	FY18	>3% growth in 1Q18
BRAC Analysis CD x°	Metastatic Breast Cancer Indication	2H FY18	FDA submission complete; expect decision in 3Q18
genesight	Additional Reimbursement	FY18	Successful prospective study
Vectra DA	ACR Guidelines & Reimbursement	2H FY18	Increased Medicare rate under PAMA
♦ Prolaris [,]	Additional Reimbursement	FY18	Increased Medicare rate under PAMA
EndoPredict®	Increased Adoption in U.S.	FY18	2% market share run rate at end of 1Q18
Path Path	Additional Reimbursement	FY18	New NCCN guidelines

Uses of Cash

Uses of Cash	Current Focus	GOAL
R&D	10% of revenue	Invest in late-stage reimbursement studies
M&A	Integration of Assurex and Sividon Acquisitions	 \$100M in potential milestones next two years Assess opportunities that fit 4in6 strategy with potential for near-term accretion
Debt Repayment	\$74M	Reduce with excess free cash flow
Share Repurchase	\$174M authorized	Opportunistic open market purchases

Worldwide Leader in Personalized Medicine

- We are entering the golden age for personalized medicine
- Molecular diagnostics are the keystone for improving patient outcomes while eliminating waste in healthcare spending
- Myriad is the pioneer of "research-based" and "education-centric" business model for molecular diagnostics
- We are the best positioned company to lead this revolution in healthcare

